Human androgen rec Human androgen rec Human immune syste TrpE/Androgen rece TrpE/N-terminal do Human immune syste Single nucleotide Single nucleotide Single nucleotide Single nucleotide Single nucleotide Phcxv-1-controlle

Rabbit progesteron PhCuV\*-1 rabbit pr Rabbit pr Mabbit pr Progesteron PhCuV\*-1 promoter sequence of a 1.23 Human progesterone duman progesterone duman progesterone

Sequence encoding Kuman breast cance TrpE/AR androgen-b cuman secreted pro

Green fluorescent Human secreted

Human androgen rec DNA encoding novel Andorgen receptor Fused androgen rec

Signal transductió Rat androgen recep Human DNA for stag

Full-length rat an

us-09-497-822c-18.rng

Sequence:

Run on:

Searched:

Database

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Human; AIB1; amplified in breast cancer 1; androgen receptor; AR; prostate cancer; chromosome X;\ ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                           /product= "Human androgen receptor (AR) protein"
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1115..3874
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Androgen receptor
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     GenCore version 5.1.4\_p5\_4578 Copyright (c) 1993 - 2003 Compugen Ltd.
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                                                                                                                                                                                                                     2185239 seqs, 1125999159 residues
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                                                    nucleic search, using sw model
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Listing first 45 summaries
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Gapop 10.0 , Gapext 1.0
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Chang C;

DNA encoding novel Rat androgen recep Human DNA for stag

Result No.

2002-206195/26

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P-PSDB; AAE19061
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The invention relates to a method for assessing the risk of prostate cancer in a human subject. The method involves determining the length of the contiguous CAG or CAA repeats in both AIBI (Amplified In Breast cancer 1) gene alleles or contiguous CAG, CAA or GGN repeats in the androgen receptor gene of the subject. The method is useful for assessing a subject s risk for acquiring or developing prostate cancer. The present sequence is human androgen receptor (AR) gene. Human AR gene is located on X chromosome. Assessing the risk of acquiring or developing prostate cancer in a human subject, comprises determining the length of the contiguous CAG, CAA and/or GGN repeats in the AIB1 gene and/or androgen receptor gene of the subject -Disclosure; Page 61-62; 86pp; English

4321 BP; 966 A; 1281 C; 1168 G; 906 T; 0 other;

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AGCTGCTAAAGACTCGGAGGAAGCAAGGAAAGTGCCTGGTAGGACTGACGGCTGCCTTTG 1046 1166 1226 1285 1345 1405 1465 657 358 418 478 537 597 986 238 298 813 873 931 Gaps 59 CCGCCCCCCCCCCCCCTCGCCCAGCGNTGNCAGNCCGAGTTTGCAGAGAGGTAACTCC CTTTGGCTGCGGGCGGGCGAGNCTAGCTGCACATTGCAAAGAAGGCTCTTAGGAG-CAGG CGACTGGGGAGCGGCTTCAGCACTGCAGCCACGACCNGCCTGGTTAGGCTGCACGCGGAG CGAGATCCCGGGGAGCCAGCTTGCTGGGAGAGCGGGAACGGTCCGGAGCAAGCCCAGAGG CAGAGGAGGCGACAGAGGGAAAAAGGGCCCNAGCTAGCCGCTCCAGTGCTGTACAGNAGC CGAA-GGACGCACCACCCAGCCCCAGCCCGGCTCCAGCGACAGCNAACGCCTCTTGCA-DB 24; Length 4321; 24; Indels 16; Query Match 82.4%; Score 4188.2; Best Local Similarity 99.1%; Pred. No. 0; Matches 4297; Conservative 0; Mismatches 1286 1346 1406 299 419 479 598 1227 538 658 754 09 874 932 179 239 1047 1107 359 1167 987 Q ò q ŏ dd QΥ g g δ qq QY Db 0.y DD QY ag QY g οy QΥ ò ద ò

2245 2305 2365 2425 2185 1945 2005 2065 1364 1424 1705 1765 1016 1825 1076 1885 1136 1196 1304 1585 836 968 926 TAGGGGGCACTTCGACCATTTCTGACAACGCCAAGGAGTTGTGTAAGGCAGTGTCGGTGT TGCTGGGCCCCACTTTCCCCGGCTTAAGCAGCTGCTCCGCTGACCTTAAAGACATCCTGA GCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGCAGGAAGCAGTATCCGAAGGCA AGCAGCAGCAGCAGCAGGGTGAGGTTCTCCCCAAGCCCCATCGTAGAGGCCCCACAG GAAGGGTCTACCCTCGGCCGCCGTCCAAGACCTACCGAGGAGCTTTCCAGAATCTGTTCC AGAGCGTGCGCGAAGTGATCCAGAACCCGGGCCCCCAGGCACCCAGAGGCCGCGAGCGCAG CACCTCCCGGCCCCAGITTGCTGCTGCTGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC GCTACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCAGTCGGCCCTGGAGTGCC 2066 2126 1365 1425 2246 1485 2306 1545 2366 1605 2426 1665 2486 2546 1197 1257 1305 2186 1946 717 1646 1706 1766 1017 1826 1077 1886 1137 2006 1526 111 1586 837 897 957 g g g g 셤 Ω ò qq qq ŏ Ω Qγ qq οy g δŽ ò δ Qγ Db g QΥ QQ δλ Ω Db g οy q δy δŏ qq Óγ q οy δ

2865 AAAGAG	3686 ATAAAT        2925 ATAAAT	3746 TGACTC        2985 TGACTC	3806 AGGCTT        3045 AGGCTT	3866 TTGAAG        3105 TTGAAG	3926 TAGTGT        3165 TAGTGT	3986 TCAATG	4046 GCTTCC         3285 GCTTCCC	4106 TCATGG7          3345 TCATGG7	4166 TCGCCCC          3405 TCGCCC	4226 GTGTCCG 	4286 TCCTGTG 	4346 AAAAATT         3585 AAAAATT	4406 GCAAAAG          3645 GCAAAAG	4466 ACTCCGT         3705 ACTCCGT	4526 CACACAT                   3765 CACACAT	4586 CCAAGAT         3825 CCAAGAT	4646 ACCCTATT 	4706 TGCACTAC          3945 TGCACTAC
qq	Qy Db	Qy Db	Qy	Qy Db	Qy Db	QY	Qy Db	Qy	QY	QY .	Qy Db	Oy Db	Qy Db	QY	QY	Qy	Qy	QY
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QQ	Qy	O <sub>Y</sub>	Qy Db	oy Q	λο qq	λ δ	Oy 2	Qy 3	0y 3	Qy 3 Db 2	Oy 3	Qy 3 Db 2	Qy 3 Db 2	Qy 3. Db 2.	Oy 3,	Oy 35 Db 27	Oy 35 Db 26	Qy 36

,~å\* 2984 3805 3865 3044 3104 PGTGCTGGACACGACAACAACCAGCCCGACTCCTTTGCAGCCTTGCTCTCTAGCC 3985 4105 GCCGCTGAAGGAAACAAAAGTACCTGTGCGCCAGCAGAAATGAT:GCACTATTG 2924 4165 3404 4285 GCATGAAAGCACTGCTACTCTTCAGCATTATTCCAGTGGATGGGCTGAAAAATC 4345 4465 4525 3704 4585 3764 3824 4645 TTTCCCCACCCCAGCTCATGCCCCTTTCAGATGTCTTGTGCCTGTTATAACTC 4705 4765 GGCTATGAATGTCAGCCCATCTTTCTGAATGTCCTGGAAGCCATT3AGCCAGGTG GCAACTTACACGTGGACGACCAGATGGCTGTCATTCAGTACTCCTGGATGGGGC TGTTTGCCATGGGCTGGCGATCCTTCACCAATGTCAACTCCAGGATGCTCTACT GAATGAGGCACCTCTCTAAGAGTTTGGATGGCTCCAAATCACCCCCCAGGAAT GAAAAAATCCCACATCCTGCTCAAGACGCTTCTACCAGCTCACCAAGCTCCTGG PGCAGCCTATTGCGAGAGAGCTGCATCAGTTCACTTTTGACCTGCTAATCAAGT g

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Proapoptotic; dependence domain; p75NTR; androgen receptor; DCC; huntingtin polypeptide; Machado-Joseph disease; SCA1; SCA2; SCA6; atrophin-1; cell death; apoptosis; Huntington's disease; head trauma; Alzheimer's disease; Kennedy's disease; spinocerebellar ataxia; stroke; dentatorubropallidoluysian atrophy; cell proliferation; cell survival; neoplastic; malignant; autoimmune; fibrotic; ss.
                                                                                                                                   This invention describes novel pure proapoptotic dependence peptides which comprise a sequence of an active dependence domain selected fro dependence polypeptides consisting of p75NTR, androgen receptor, DCC,
                                                                                                                         TCTTATGCCACGGGAAGTTTAGAGAGCTAAGATTATCTGGGGAAATCAAAACAAAAAAA
TCTTCCCTCCCTATCTAACCCTCCCATGGCACCTTCAGACTTTGCTTCCCATTGTGGCTC
                                                                                 CTATCTGTGTTTTGAATGGTGTTGTATGCCTTTAAATCTGTGATGATCCTCATATGGGCCC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     to develop products
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          "androgen receptor"
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     proapoptotic dependence peptides, used
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532..3288
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huntingtin polypeptide, Machado-Joseph disease gene product, SCA1, SCA2, SCA6 and atrophin-1 polypeptide. The proapoptotic peptides are capable of inducing cell death and can be used to develop products to mediate or inhibit apoptosis. The methods can be used for reducing tithe severity of a proapoptotic dependence domain mediated pathological conditions e.g. Huntington's disease, Alzheimer's disease, Kennedy's disease, Appincertebellar ataxias, dentatorubropallidoluysian atrophy. Machado-Joseph disease, stroke or head trauma. They can also be used for reducing the severity of a pathological condition mediated by upregulated each proliferation or cell survival e.g. neoplastic, malignant, autoimmune or fibrotic conditions. This sequence encodes a human androgen receptor described in the method of the invention.
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                                                                                                                                                            Sequence 3715 BP; 841 A; 1055 C; 1001 G; 818 T; 0 other;
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Qy	Qy	Qy	Qy	Qy Dp	Qy	Q <sub>Y</sub>	Qy Db	O <sub>Y</sub>	Qy	Qy	oy do	Qy	ογ Dp	7 da	Oy 4	Oy 4 Db 2	Qy 4 Db 2
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759 TAGCCCCAGGCAGCAGCAGCAGCAGGAGGTGAGGATGTTCTCCCCAAAGCCCATCGTAG 818 2175 AGGCCCCACAGGTACCTGGATGAGAACAGCAACCTTCACAGCCGCAGTCGGC 2234 11111111111111111111111111111111111	22	23.9	24 24 10	247,	253		265	27	~ ~	283	5 CGAGAGCCTAGGCTGCTCTGGCAGCGCTGCAGCAGGAGCTCCGGGACACTTGAACTGCC 289	5 GTCTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGAGCTGCGTACCAGAGTCG 295.	301	307	5 GGCGGCGCAGTGCCGCTATGGGGACCTGGCGAGCCTGCATGGCGCGGGTGCAGCGGGACC 313.	5 CGGTTCTGGGTCACCTCAGCCGCCGCTTCCTCATCCTGGCACACTCTCTTCACAGCCGA 319	195 AGAAGCCCAGTTGTATGGACCGTGTGGTGGTGGGGGGGGG
Oy Db	Oy Dp	Oy Db	ζ, QΩ	Qy 2	OY 2 Db 1	0y 2 Db 1	Oy 2 Db 1	Oy 2 Db 1	0y 2 Db 1	Oy 2'	Oy 283. Db 147	Oy 289	Qy 29	Qy 30 Db 16	Qy 307 Db 171	Qy 313 Db 177	Qy 31 Db 18

CGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGGCGAGGCGGAGCTGT CCCCTACGGCTACACTCGGCCCCTCAGGGGCTGGCGGGCCAGGAAAGCGACTTCAC PGTCAAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGGACAT | TTGGAGACTGCCAGGGACCATGTTTGCCCATTGACTATTACTTTCCACCCAGAA TGCCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACATGTGG TGCAAGGTCTTCTTCAAAAGAGCCGCTGAAGGGAAACAGAAGTACCTGTGCGCCAG AATGATTGCACTATTGATAAATTCCGAAGGAAAATTGTCCATCTTG;CGTCTTCG CTACAGGAGGAAGGAGGGTTCCAGCACCACCAGCCCCACTGAGGAGAACCCA CTGACAGTGTCACACATTGAAGGCTATGAATGTCAGCCCATCTTTCTGAATGTCCT SCCITGCICTAGCCICAAIGAACIGGGAGAGAGACAGCITGIACACGIGGICAA CCAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGCACAA TCACCCCCAGGAATTCCTGTCCATGAAAGCACTGCTACTCTTCAGCATATTCC genes in

Diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver tumor in a patient, involves detecting the level of expression of two or mora genes

liver tissue sample

Vockley JG;

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Peres-Da-Silva

02-OCT-2000; 2000US-237054P. 02-OCT-2001; 2001WO-US30589

(GENE-) GENE LOGIC INC Horne D, Alvares C,

WPI; 2002-426119/45.

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Gene; liver cancer; ds; hepatocellular carcinoma; hepatotropic; metastatic liver tumour; cytostatic; expression profile; disease state; disease progression; drug toxicity; drug efficacy; drug metabolism.
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The invention relates to a novel method for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma or metastatic liver cancer from tumour in a patient, and differentiating metastatic liver cancer from chaptocellular carcinoma in a patient, involving detecting the level of expression of two or more genes represented in ABN3503-ABNS7455 in a expression of two or more genes represented in ABN3503-ABNS7455 in a carcinoma sample. The method is useful for diagnosing and detecting cytostatic activity. The method is useful for diagnosing and detecting the progression of liver cancer, hepatocellular carcinoma and metastatic liver carcinoma in a patient. The method is useful for identifying expression profiles which serve as useful diagnostic markers as well as markers that can be used to monitor disease states, disease progression, drug toxicity, drug efficacy and drug metabolism.

Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic format directly from WIPO cat fip. wipo.int/pub/published_pct_sequences.
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                                                                                                                                                                                                                                                                                                                                                                                                                  Sequence 3715 BP; 841 A; 1055 C; 1001 G; 818 T; 0 other;
                                                                                                                                                                                    Claim 1; SEQ ID NO 2300; 298pp; English.
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//roduct= full-length 918 residue hAR
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                                                                                                                                                                                   protein; steroid hormone; ss.
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iive 0; Mismatches
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                                                                AAQ12001 standard; cDNA; 3715
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Db 3508 TTTGCTTCCCATTGTGGCTCCTATCTGTG OY 4926 TGATGATCCTCATAGGCCCAGTGTCAAC Db 3568 TGATGATCCTCATAGGCCCAGTGTCAAC OY 4986 GCCACAAACGTTACTTATCTTATCT Db 3628 GCCACACAAACGTTACTTATTATGCC OY 5046 GGAAATCAAAACAAAAA 5062 Db 3688 GGAAATCAAAAAAA 3704	RESULT 5 AAT63407 ID AAT63407 standard; cDNA; 3569 BP. XX AC AAT63407; XX NDT 22-JUN-1997 (first entry) XX XX	Androgen receptor cDNA.  XX  Androgen receptor; acidic fibroblast growth factor; aFGF;  KW  Antisense; benign prostatic hyperplasia; prostate cancer;  KW  ds.  XX  NX  NX  XX  XX  XX	Key CDS misc_feature misc_feature	misc_feature	XX XX XX XX XX XX 20-SEP-1996; 96WO-US15081. PX 20-SEP-1995; 95US-0004018.	XX PA (WORC-) WORCESTER FOUND BIOMEDICAL RES XX PI Zamecnik PA; XX XX XX PI 1997-202879/18. DR PPSDB: AAW14783.	antisense inhibit perplasis ; 51pp; E 407) code s of sele cells in
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GAAACTACAGGAGGAAGGAGGCTTCCAGCACCACCACCACTGAGGAGACAACCCA         3845	GTGGGCCAAGGCCTTGCCTGGCTTCCGCAACTTACACGTGGACGACCACAGATGGCTGTCAT 4085 GTGGGCCAAGGCCTTGCCTTCCGCAACTTACACGTGGACGACACAGATGGCTGTCAT 2738 GTGGGCCAAGGCCTTGCCTTCGCCAACTTACACGTGGACGACACAGATGGCTGTCAT 2738 TCAGTACTCCTGGATGGGGCTCATGGTGTTTGCCATGGGCTGGCGATCCTTCACCAATGT 4145 T1111111111111111111111111111111111	CAACTCCAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGCACAA 4205	CCAAATCACCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTCTTCAGCATTATTCC 4325	ACTCGATCGTATCATTGCATGCAAAAAAAAACCCACATCCTGCTCAAGACGCTTCTA 4445	TTTGACCTGCTAATCAAGTCACATGGTGAGCGTGGACTTTCCGGAAATGATGGCAGA 4565	CACCCAGTGAAGCATTGGAAACCCTATTCCCCACCCCAGCTCATGCCCCTTTCAGATG 4685 CACCCAGTGAAGCATTGGAAACCCTATTTCCCCACCCCA	

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receptor sequence or antisense oligonucleotides (see also AAT63406) to the human acidic fibroblast growth factor gene (see also AAT63197-99). The methods are esp. useful for the treatment of benign prostatic hyperplasia and prostate cancer.
                                                                                                                                                                    TCCCGCAGGTGGGCAGCTAGCTGCAGCGACTACCGCATCATCACAGCCTGTTGAACTCTT
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                                         BP; 796 A; 1009 C; 974 G; 790 T;
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0; Mismatches
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larity 99.7%; Pred. No. 0;
Conservative 0; Mismatche
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Matches 3569; Conserv
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2581 2761 2701 1428 3121 3241 3181 948 1908 3481 3541 2028 CCGCTGACCTTAAAGACATCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGC CCACTTCCTCCAAGGACAATTACTTAGGGGGCACTTCGACCATTTCTGACAACGCCAAGG **AGTTGTGTAAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCA.** TCTGA GTCCAGGGGAACAGCTTCGGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCCG 1129 GTCCAGGGGAACAGCTTCGGGGGATTGCATGTACGCCCACTTTTGGGACTTCCACCC CTGTGCGTCCCACTCCTTGTGCCCCATTGGCCGAATGCAAAGGTTCTCTGCTAGACGACA GCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTfACACCA AAGGGCTAGAAGGCGAGAGCCTAGGTGCTGGCAGCGCTGCAGCAGGAGGTCCGGGA CACTTGAACTGCCGTCTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTG CGCCGCCTCCCCATCCCCACGCTCGCATCAAGCTGGAGAACCCGCTGGACTACGGCAGCG CGTACCAGAGTCGCGACTACTACAACTTTCCACTGGCTCTGGCCGGACCGCCCCCCTC CCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGGGACCTGGCGAGCCTGCATGGCGCGG GTGCAGCGGGACCCGGTTCTGGGTCACCCTCAGCCGCCGCTTCCTCATCCTGGCACACTC TCTTCACAGCCGAAGAAGGCCAGTTGTATGGACCGTGGTGGTGGTGGTGGTGGTGGTGGTGGC CTGTAGCCCCCTACGGCTACACTCGGCCCCCTCAGGGGCTGGCGGGCCAGGAAAGCGACT TCACCGCACCTGATGTGTGTACCCTGGCGGCATGGTGAGCAGGTGCCCTATCCCAGTC CCACTTGTGTCAAAAGCGAAATGGGCCCCTGGATGGATAGCTACTCCGGACCTTACGGGG ACATGCGTTTGGAGACTGCCAGGGACCATGTTTTGCCCATTGACTATTACTTTCCACCC 889 2462 949 2522 1009 2582 1069 2642 2702 2822 2762 2882 1369 2942 1489 3062 1549 1429 3002 3122 1609 3182 1669 3242 1729 3302 3362 1849 1789 3422 1909 3482 1969 ŏ Q g Ω ò Ωp δ Op õ Dp οy Op QQ οy ρý Db Ω Db δy OD οy QQ οy qq QΥ g Qγ g δ g g ōλ δy g ò a

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Complementary DNA sequences derived from the CDNA may be used as probes
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                                       DNA encoding androgen receptor protein – useful for transforming eukaryotic hosts for protein expression and subsequent antibody protein expression and subsequent and
                                                                                                    4682 GAIGICIICIGCCIGITATAACICIGCACIACICCICCIGCAGIGCCIIGGGGAAITICCI
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                         Location/Qualifiers 363..3122
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                AAN91772 standard; cDNA; 3569 BP.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     Human androgen receptor cDNA.
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P-PSDB; AAP93109.
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detect the presence of androgen receptor (AR mRNA in tumour cells, and detect AR gene defects using DNA hybrisidation assays. Sequence 3569 BP; 796 A; 1008 C; 975 G; 790 T; 0 other;	ery Match st Local Similarity 99.6%; Pred. No. 0; tches 3568; Conservative 0; Mismatches 1; In	TACTICAGIGGACACIGAATTIGGAAGGIGGAGGA 1	1562 ITTIGITITITICITITAAGAICIGGGCAICITITGAAICIACCCTICAAGIAITAAGAG 1621 	18	CTGGAGCT 17 	1742 TCCCGCAGGTGGGCAGCTAGCTGCAGCGCATCATCACAGCCTGTTGAACTCTT 1801 	1802 CTGAGCAAGAGAAGGGGGGGGGTAAGGGAAGTAGGTGGAAGATTCAGCCAAGCTCAA 1861 	1862 GGATGGAAGTGCAGTTAGGGCTGGGAAGGGTCTACCCTCGGCCGCCGTCCAAGACCTACC 1921 	1922 GAGGAGCTTTCCAGAATCTGTTCCAGAGCGTGCGCGAAGTGATCCAGAACCGGGCCCCA 1981 	1982 GGCACCAGAGGCCGCGAGCGCAGCACCACCTCCCGGCGCCAGTTTGCTGCTGCTGCAGCAGC 2041	2042         AGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCA	2102 AGCAGCAAGAGACTAGCCCCAGGCAGCAGCAGCAGGAGGGGGGGG	2162 AAGCCCATCGTAGAGGCCCCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCAC 2221	2222 AGCGGCAGTCGGCCCTGGAGTGCCACCCCGAGAGGTTGCGTCCCAGAGCCTGGAGCG 2281	2282 CCGTGGCCGCCAGCAGGGGCTGCCGCAGCTGCCAGCACCTCCGGACGAGGATGACT 2341 	2342 CAGCTGCCCCATCCACGTTGTCCCTGCTGGCCCCACTTTCCCCGGCTTAAGCAGCTGCT 2401 	2402 CCGCTGACCTTAAAGACATCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGC 2461 
S X X &	Ma. Ma.	Qy	Oy Dp	Qy	oy Db	Qy Db	Qy	Qy Dp	Q <sub>Y</sub>	Qy Db	oy Db	Qy Db	Oy Dp	Qy	Qy Db	Oy Dp	Oy

2581 1068 2641 2521 2881 2941 1428 3061 3121 1608 3181 1668 3241 1728 3301 3361 1848 3421 1548 3481 1968 3541 2028 3542 AGAAGACCTGCTGATCTGTGGAGATGAAGCTTCTGGGTGTCACTATGGAGCTCTCACAT 3601 AGTIGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCATCTGA CCACTICCTCCAAGGACAATTACTTAGGGGGCACTTCGACCATTTCTGACAACGCCAAGG GCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTTACACCA AAGGGCTAGAAGGCGAGAGCCTAGGCTGCTCTGGCAGCGCTGCAGCAGGAGCTCCGGGA CACTTGAACTGCCGTCTACCCTGTCTCTTACAAGTCCGGAGCACTGGACGAGGCAGCTG CGCCGCCTCCCCATCCCCACGCTCGCATCAAGCTGGAGAACCCGCTGGACTACGCCAGCG CCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGGACCTGGCGAGCCTGCATGGCGCGG CTGTAGCCCCCTACGGCTACACTCGGCCCCCTCAGGGGCTGGCGGCCCAGGAAAGCGACT TCACCGCACCTGATGTGTGTACCCTGGCGGCATGGTGAGCAGAGTGCCCTATCCCAGTC ACATGCGTTTGGAGACTGCCAGGGACCATGTTTTGCCCATTGACTATTACTTTCCACCCC 2522 2462 1009 2582 1069 2702 1189 1249 1309 1369 ( 2762 2822 2882 3002 1549 1489 3062 3122 1609 3182 1669 3242 1729 3302 1789 3362 1849 3482 1969 qq δy Ω g δ Q Óλ qq Ω Db δ Dβ QY Db δy qq Oy Oy QQ Οÿ QQ qq qq ŏλ δλ Óλ q δŽ qq Óγ g δy g δ g ò

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	602	3662 CCAGCAGAAATGATTGCACTATTGATAAATTCCGAAGGAAAAATTGTCCATCTTGTCGTC 	3722 TTCGGAAATGTTATGAAGCAGGGATGACTCTGGGAGCCCGGAAGCTGAAGAAACTTGGTA 	3782 ATCTGAAACTACAGGAGGAAGGAGGCTTCCAGCACCACCAGCCCCACTGAGGAGACAA	3842 CCCAGAAGCTGACAGTGTCACATTGAAGGCTATGAATGTCAGCCCATCTTTCTGAATG	3902 TCCTGGAAGCCATTGAGCGGGGTAGTGTGTGTGGGACACGACAACAACCAGCCCGACT 	3962 CCTTTGCAGCCTTGCTCTAGCCTCAATGAACTGGGAGGAGACACTTGTACACGTGG 	022 TCAAGTGGGCCAAGGC 	4082 TCATTCAGTACTCCTGGATGGGCTCATGGTGTTTGCCATGGCTGGC	4142 ATGTCAACTCCAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGC	4202 ACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGCACCTCTCAAGAGTTTGGAT 	4262 GGCTCCAAATCACCCCCAGGAATTCCTGTGCATGAAAGCACTGCTACTCTTCAGCATTA 	4322 TICCAGIGGAIGGGCIGAAAAATCAAAAAITCITIGAIGAACTICGAAIGAACTACATCA 	4382 AGGAACTCGATCGTATCATTGCATGCAAAAGAAAAATCCCACATCCTGCTCAAGACGCT 	4442 TCTACCAGCTCACCAAGCTCCTGGACTCCGTGCAGCCTATTGCGAGAGAGCTGCATCAGT [	4502 TCACTTTTGACCTGCTAATCAAGTCACACATGGTGACGGTGGACTTTCCGGAAATGATGG 	4562 CAGAGATCATCTCTGTGCAAGTGCCCAAGATCCTTTCTGGGAAAGTCAAGCCCATCTATT 	4622 TCCACACCCAGTGAAGCATTGGAAACCCTATTTCCCCACCCCAGCTCATGCCCCCTTTCA
qq	Qy Db	Qy	Oy Db	Oy Db	Qy	Qy	ζζ	Qy Dp	Qy Db	ζ QΩ	ζ Q	Qy	OY. Ob	QY	Oy Db	Q P	oy Op	Qy

chromosome 3408 3468 5041 3168 3228 4801 4921 4981 3288 The invention relates to isolated polynucleotide (I) and polypeptide (II) sequences. (I) is useful as hybridisation probes, polymerase chain reaction (PCR) primers, oligomers, and for chromo and gene mapping, and in recombinant production of (II). The SS seful in mutations 3469 GCCAGCCACACAAAGTTTACTTATCTTATGCCACGGGAAGTTTAGAGAGCTAAGATTAT 4922 TCTGTGATGATCCTCATATGGCCCAGTGTCAAGTTGTGCCTTGTTAACAGCACTACTGT GCCAGCCACACAAACGTTTACTTATCTTATGCCACGGGAAGTTTAGAGAGCTAAGATTAT 3409 TCTGTGATGATCCTCATATGGCCCAGTGTCAAGTTGTGCTTGTTTACAGCACTACTGT 3109 TCCACACCCAGTGAAGCATTGGAAACCCTATTTCCCCACCCCAGCTCATGCCCCTTTCA CTAITGAIGTACAGICIGICAIGAACAIGIICCIGAAIICIAIITIGCIGGGCIITIIIII New isolated polynucleotide and encoded polypeptides, useful in diagnostics, forensics, gene mapping, identification of mutation responsible for genetic disorders or other traits and to assess biodiversity Human; chromosome mapping; gene mapping; gene therapy; forensic; food supplement; medical imaging; diagnostic; genetic disorder; DNA encoding novel human diagnostic protein #29276 Claim 1; SEQ ID No 29276; 103pp; English. BP. RESULT 7 AAS93472/c ID AAS93472 standard; cDNA; 3590 Tang YT; 31-MAR-2000; 2000US-0540217. 23-AUG-2000; 2000US-0649167. 30-MAR-2001; 2001WO-US08631 (first entry) Drmanac RT, Liu C, WPI; 2001-639362/73. P-PSDB; ABG29285. (HYSE-) HYSEQ INC. WO200175067-A2. Homo sapiens 13-FEB-2002 11-0CT-2001. AAS93472; 3529 3289 4982 4742 3229 4802 δy q δλ QQ ŏ qq OD q δ ф Óγ

polynucleotides are also used in diagnostics as expressed sequence tags for identifying expressed genes. (I) is useful in gene therapy techniques to restore normal activity of (II) or to treat disease states involving (II). (II) is useful for generating antibodies against it, detecting or consolidating a polypeptide in tissue, as molecular weight markers and as a food supplement. (II) and its binding partners are useful in medical consolidates involving aberrant protein expression or biological activity. The polypeptide and polynucleotide sequences have applications in diagnostics, forensics, gene mapping, identification of mutations in consolidation of produce other types of data and products dependent on DNA and and adagnostic coding sequences of the invention.

Conditional sequence data for this patent did not appear in the printed sequence data for this patent did not appear in the printed sequence consolidation, but was obtained in electronic format directly from WIPO ftp.wipo.int/pub/published\_pct\_sequences \$

Sequence 3590 BP; 807 A; 977 C; 1016 G; 790 T; 0 other;

1621 3471 1681 3411 1741 3531 1801 3350 TCCCGCAGGTGGGCAGCTAGCTGCAGCGACTCATCATCACAGCCTGTTGAACTCTT 3291 CTGAGCAAGAAGAGGGGAGGCGGGGTAAGGGAAGTAGGTGGAAGATTCAGCCAAGCTCAA 1861 1921 2041 3052 2874 GAGGAGCTTTCCAGAATCTGTTCCAGAGCGTGCGCGAAGTGATCCAGAACCCGGGGCCCCA 1981 2101 2994 2161 2934 2222 AGCCGCAGTCGGCCCTGGAGTGCCACCCCGAGAGAGGTTGCGTCCCAGAGCCTGGAGCCG 2281 Gaps TAATAACTCAGTTCTTATTTGCACCTACTTCAGTGGACACTGAATTTGGAAGGTGGAGGA TITIGITITITICITITIAAGAICTGGGCATCTTTTGAATCTACCCTTCAAGTATTAAGAG ACAGACTGTGAGCCTAGCAGGGCAGATCTTGTCCACCGTGTGTCTTCTTCTGCACGAGAC 1742 TCCCGCAGGTGGGCAGCTAGCTGCAGCGACTACCGCATCATCACAGCCTGTTGAACTCTT AAGCCCATCGTAGAGGCCCCAACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCAC GGCACCCAGAGGCCGCGAGCGCACCTCCCGGCGCCAGTTTGCTGCTGCTGCAGC AGCAGCAAGAGACTAGCCCCAGGCAGCAGCAGCAGCAGGAGGTGAGGATGGTTCTCCCC DB 23; Length 3590; 25; 3; Indels Score 3498.2; Pred. No. 0; 0; Mismatches Query Match
Best Local Similarity 99.2%;
Matches 3565; Conservative Query Match 1502 1622 3470 1562 1802 3290 3230 1922 3170 3110 3051 1862 1982 2042 2102 2993 2162 ο̈́λ q οy q δ Ω δ Q οŽ a Q δy Qγ g οqα οχ δ q Ω qq δy g Ω a

2814 2401 2461 2761 2881 2941 2154 3001 3061 3181 1974 1914 3241 3361 CCGTGGCCGCCAGCAAGGGGCTGCCGCAGCTGCCAGCACCTCCGGACGAGGATGACT CCGCTGACCTTAAAGACATCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGC AGCAGGAAGCAGTATCCGAAGGCAGCAGCAGGGAGGGGAGGCCTCGCGGGCTC AGTTGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCATCTGA CTGTGCGTCCCACTCCTTGTGCCCCATTGGCCGAATGCAAAGGTTCTCTGCTAGACGACA CCACTTCCTCCAAGGACAATTACTTAGGGGGCACTTCGACCATTTCTGACAAGGG GCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTTACACCA CACTTGAACTGCCGTCTACCCTGTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTG CCTGGGCGGCTGCGGCGCGCAGTGCCGCTATGGGGACCTGGCGAGCCTGCATGGCGCGG TCTTCACAGCCGAAGAAGGCCAGTTGTATGGACCGTGTGGTGGTGGTGGTGGTGGTGGTGGTGGT CGTACCAGAGTCGCGACTACTACAACTTTCCACTGGCTCTGGCCGGACCGCCGCCCTC CTGTAGCCCCCTACGCCTACACTCGGCCCCCTCAGGGGCTGGCGGGCCAGGAAAGCGACT 2813 2753 2873 2342 2402 2693 2462 2573 2633 2513 2762 2522 2642 2453 2702 2393 2333 2822 2273 2882 2213 2153 2093 2942 3002 3062 2033 1973 3182 3122 1913 3242 1853 3302 δ Db Qγ ρp ōλ g qq Ω Dp δ QY Db QQ g QY δλ οy QQ οy Pp QΥ g Qy g ŏ g ŏ qq ò a q ÓΥ ò qq ò

1932 CTGTRAGCCCCTACGGCTACACCTGGGCGCTGGGGGCTGGGGGCGGGGCGGGC

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                                                                           CCCAGCTCATGCCCCTTTCAGATGTCTTCTGCCTGTTATAACTCTGCACTACTCCTCTG 4720
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                                      4541 TGGACTTTCCGGAAATGATGGCAGAGATCATCTGTGCAAGTGCCCAAGATCCTTTCTG 4600
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                                                                                              113 TIGITTACAGCACACACCACACACACACTAACTTATGCACGGG
                                           Rat androgen receptor; monoclonal antibody; ployclonal antibody;
                                                                                                                                                                                             Location/Qualifiers
996..3702
/*tag= a
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                                                                                                                                                                                                                                AAN91773 standard; cDNA; 4180
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                                                                                                                                                                                                                                                              Rat androgen receptor cDNA.
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                                                                                                                                                                                                                                                                             cancer; probe.
                                                                                                                                                                                                                                                                                      Rattus rattus,
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ACCGCATCATCACAGCCTGTTGAACTCTTCTGAGCAAGAGAAGGGGAGGCGGGGTAAGGG
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                                                                            1653 TCCACCGTGTGTCTTCTGCACGAGACTTTGAGGCTGTCAGAGCGCTTTTTGCGTGGT
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                                                                                                                                                                             758 ATCCCGGGGAGCCAGCTTGCTGGAAGGGGGAACGGTCCGGAGCAAGCCCAGAGGCAGA
                                                                                                                                                                                      -GGACGCACCACCCAGCCCCAGCCCGGCTCCAGCGACAGCNAACGCCTCTTGCANGCGT
                                                                                                                                                                                                                                                     TCGAAGCCGCCCC-GGAGCTGCCCTTTCCTTCGGTGAAGTTTTTAAAAGCTGCTAA
                                                                                                                                                                                                                                                                                                                    297 CTCCTACCC-------CTACCCTCTGGGTCCCTCTGGGG
                                                                                                                                                                                                                                                                                                                                                                         1116 TGCCTCAGTCGGCTACTCTCAGCCAACCCCCTCACCACCCTTCTCCCCACCCGCCCCC
                                                                                                                                                                                                                                                                                                                                                                                          336 GGACTAGGCAGGCTTC-----CTGGCCAGCC
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          transforming
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    DNA encoding androgen receptor protein – useful for transforming eukaryotic hosts for protein expression and subsequent antibody
                                                                                                                                                  Length 4180;
                                                                                                                                                                   306;
                                                                                                    nseq
                                                                                              Complementary DNA sequences derived from the cDNA may be used detect the presence of androgen receptor (AR) mRNA in tumour to detect AR gene defects using DNA hybrisidation assays.
                                                                                                                            Sequence 4180 BP; 1024 A; 1149 C; 1083 G; 924 T; 0 other;
                                                                                                                                                                664; Indels
                                                                                                                                                 Score 2486.4; DB 10;
                                                                                                                                                        Pred. No. 0;
0; Mismatches
                  Joseph DR, Lubahn
  NORTH CAROLINA
                                                                                 English
                                                                                                                                              Query Match
48.9%;
Best Local Similarity 77.9%;
Matches 3416; Conservative 0
                                                                             Disclosure; Fig. 5; 41pp;
                  EW,
UNIVERSITY OF
                                WPI; 1989-324206/44.
P-PSDB; AAP93110.
                 Wilson
                 French FS,
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2562   CATTCTGACAACGCCAAGGAGTTGTGTAAGGCAGTGTCGGTGTCCATGGGCCTGGGTGT	734 GGAGCACTGGAACATCTGAGTCCAGGGGAGCAGCTTCGGGGCGACTGCATGTACGCGT 682 ACTITTGGGAGTTCCACCGCTGCGTCCCACTCCTTGTGCCCATTGGCCGAATGCA	AGTTCTCTGCTAGACGACGCCAGGCAAGACCCTGAAGATA	802 TTTCAAGGGAGGTTACACCAAAGGGCTAGAAG 	862 TGCAGCAGGGAGCTCCGGGACACTTGAACTGCCGTCTACCCTGTCTCTCT	22 AGCACTGGACGAGG               34 AGCAGTAGAGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGG	2000	042 CCCG        154 CCCG	3102 GGCGAGCCTGCATGGCGGGGGGCACCCGGTTCTGGGTCACCCTCAGCCGCCGCCCCTCAGCCGCCGCCCCCCGCCCCCCCC		222 TGGTGGTGGGGGGTGGTGGCGGCGGCGGCGGCGGCGGCGG	GCGGCGAGGCGGAGCTGTACGCCCTACGGCTACACCTCGGCCCCCTCAGGGCGCCCTCAGGGCGCTACACCTCGGCCCCTCAGGGGCCCTACGGCGCTACACCTCGGCCCCTCAGGGGCCCTACGGCCCTACGGCGCCCTCAGGGGCCCTACGGCGCTACACACAC	349 TAGCCCAAGCGATGCTGGGCCTGTAGCCCCCTATGGCTACACTCGGCCCCCTCAGG	3342 GGCGGCCAGGAAAGCGACTTCACCGCACCTGATGTGTGTG	3402 CAGAGTGCCCTATCCCAGTCCCACTTGTGTCAAAAGGGAAATGGGCCCCTGGATGGA	3462 CTACTCCGGACCTTACGGGGACATGCGTTTGGAGACTGCCAGGGACCATGTTTTGCCCAT	3522	3582 TCACTAT	3642 GAAACAGAAGTACCTGTGCGCCAGCAGAAATGATTGCACTATTGATAATTCCGA
Oy Db	oy oy	dy dy	QY	QY	Qy Db	QY Db	Qy Db	Qy Db	QQ Dp	οy	Db Qy	QQ	Qy Db	QY Db	Q Q	S da	Oy	QY

4719 4778 3428 4421 4481 3668 3728 3787 4181 3248 4241 3368 4361 3488 3548 4541 3608 4601 4661 4061 3188 4301 3068 2768 3761 2888 3881 2948 3941 3008 4001 3128 4121 3821 GCAGTGCCTT-GGGGAATTTCCTCTATTGATGTACAGTCTGTCATGAACATGTTCCTGAA TTTCAATGAGTACCGCATGCACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGCA CACATCCTGCTCAAGACGCTTCTACCAGCTCACCAAGCTCCTGGACTCCGTGCAGCCTAT AAATTGTCCATCTTGTCGTCTTCGGAAATGTTATGAAGCAGGGATGACTCTGGGAGCCCG CAGCCCACTGAGGAGACACCCAGAAGCTGACAGTGTCACACATTGAAGGCTATGAATG CGACAACAACCAGCCCGACTCCTTTGCAGCCTTGCTCTCTAGCCTCAATGAACTGGGGAAACTGGAGAACTGGAGACCAGCTGATTCCTTTGCTGCTGCTGTTATCTAGTCTCAAGAGCTTGGCGGA CGTGGACGACCAGATGGCTGTCATTCAGTACTCCTGGATGGGGCTCATGGTGTTTGCCAT GGGCTGGCGATCCTTCACCAATGTCAACTCCAAGATGCTCTACTTCGCCCCTGATCTGGT 4542 3609 4602 3669 4662 3729 4720 4422 3129 3189 3249 4242 3309 4302 3369 3489 4122 4182 2709 3882 2949 3942 4002 3069 4062 2769 2829 3822 2889 3009 3702 3762 Db g δy g ò g ò g Qγ g QΫ́ g Q g QY δy qq Qγ g δ qq . O. οy g Db δy PP QΥ qq pp Ω ΩŸ δ δý Qγ

methylation states of the CpG dinucleotides of (I). The array is useful clar determining genetic and/or epigenetic parameters, classification, claiming, grading, staging, treatment and/or diagnosis of astrocytomas by analysing cytosine baserotytomas, or the predisposition to astrocytomas by analysing cytosine methylations, involves obtaining a biological sample containing genomic DNA, converting cytosine bases which are denomic DNA sample, to uracil or enother base which is dissimilar to cytosine in terms of hybridisation behaviour, by chemical treatment and amplifying chemically pre-treated amplificates carry a detectable label. The method further involves of amplificates carry a detectable label. The method further involves concounting methylation status of one or more cytosine positions, and analysing methylation status of the cytosine positions by reference to bisulphite, hydrogen sulphite or disulphite. The amplification status of the cytosine positions by reference to bisulphite, hydrogen sulphite or disulphite. The amplification status of barticular interest in astrocytoma or step amplifies DNA which is of particular interest in astrocytoma or tissues, as opposed to background DNA. The amplification status of brain tissue, based on the specific genomic methylation status of brain tissue, based on the specific genomic methylation status of brain tissue, and detected in a mass spectrometer. The fragments of the amplificates are detachable molecule fragments having a typical mass which are detected in a mass spectrometer. The fragments of the amplificates are detachable series of the amplificates are detected by matrix assisted laser describulation in the mass spectrometer. Conspirity of the amplificates are detected by matrix assisted laser describulation in the mass spectrometer. The molecule is assisted laser describulation in the mass spectrometer (MADI) or using electron spray mass spectrometry (ESI). The semilation.

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Human; ds; astrocytoma; cytostatic; staging; cysteine methylation; CpG;
bisulphite; brain tissue; MALDI; ESI; electron spray mass spectrometry;
matrix assisted laser desorption/ionization mass spectrometry.
                         4839 TCTAACCCICCCAIGGCACCIICAGACIIIGCIICCCAIIGIGGCICCIAICIGIGIIII 4898
                                                      GCTTGTTTACAGCAC - TACTCTGTGCCAGCCACACACATTACTTATCTTATGCCAC 5016
                                                                                                                                       GGGAAGTITAGAGAGCTAAGAITATCTGGGGAAATCAAAAACAAAAAACAAGCAAACAAAA 5076
3788 GCATTGCCTTGGGGGAAATTCCTCTACTGATGTACAGTCTGTCATGAACATGTTCCCCAA 3847
                                                                                4958
                                                                                                                 GAATGGTGTTGTATGCCTTTAAATCTGTGATGATCCTCATATGGCCCAGTGTCAAGTTGT
                                                                                   Human DNA for staging of Astrocytomas, complement, #50.
                                                                                                                                                                                                                        ABK34013 standard; DNA; 3715 BP.
                                                                                                                                                                                                                                                                                                                                                               02-JUL-2001; 2001WO-EP07538.
                                                                                                                                                                                                                                                                                                                                                                                30-JUN-2000; 2000DE-1032529.
01-SEP-2000; 2000DE-1043826.
                                                                                                                                                                                                                                                        (first entry)
                                                                                                                                                                                                                                                                                                                                                                                                      (EPIG-) EPIGENOMICS AG.
                                                                                                                                                                      AAAAA 5082
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                                                                                                                                                                                                                                                                                                                                 WO200202808-A2.
                                                                                                                                                                                                                                                                                                                    Homo sapiens.
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Note: The sequence data for this patent did not form part of the printed specification, but was obtained in electronic

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1701 ITTTTGCGTGGTTGCTCCCGCAAGTTTCCTTCTGGAGCTTCCCGCAGGTGGGCAGCTA 1760
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                                                                                                                                                                                                                                                                    45; Gaps
                                                                                                                                                                                                                                                                                                          GAGTGCGGAGCCAGAGATCAAAAGATGAAAAGGCAGTCCAGGTCTTCAGTAGCCAAAAAAC
                                                                                                                                                                                                                                                                                                                  1521 TGCACCTACTTCAGTGGACACTGAATTTGGAAGGTGGAGGATTTTGTTTTTTTAA
                                                                                                                                                                                                                                                                                                                                                                           1641 GGGCAGATCTTGTCCACCGTGTGTCTTCTTCTGCACGAGACTTTGAGGCTGTCAGAGCGC
                                                                                                                                                                                                                                                                                                                                                                                                                                  Score 2244.2; DB 24; Length 3715;
Pred. No. 0:
                                                                                                                                                                                                                                                           Pred. No. 0;
0; Mismatches 848; Indels
                                                                                                                                                                                                                                    Sequence 3715 BP; 818 A; 149 C; 1055 G; 1693 T; 0 other;
                                                                                                                                                                                                                     ftp.wipo.int/pub/published_pct_sequences.
                                                                                                                                                                                                                                                    44.28;
76.18;
                                                                                                                                                                                                                 directly from WIPO at
                                                                                                                                                                                                                                                                 Conservative
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Matches 2838; Conserv
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The invention relates to a nucleic acid comprising a sequence (I) of at least 18 bases in length of a segment of chemically pre-treated genomic DNA which has any one of the sequences of (ABK33919-ABK34032) or its complement. Also included are an Oligonucleotide or peptide nucleic acid (or set thereof) of at least 9 nucleotides which hybridises to (I), probes for detecting cytosine methylation or singlenucleotide polymorphisms (SNP) in (I), an array of oligomers or peptide nucleic acids for analysing diseases associated with the

Novel chemically modified genomic DNA sequences, useful in the characterisation, classification, differentiation, grading, staging, treatment and/or diagnosis of astrocytomas or predisposition to

Berlin K;

Olek A, Piepenbrock C,

WPI; 2002-171649/22

Claim 1; SEQ ID No 100; 37pp; English.

astrocytomas

3168 1940 3108 2000 3048 2060 3001 2120 2952 2180 2892	2240 2832 2300 2772 2772 2360 2712 2712 2712	593 593 593 593 593 593 593 593 593 593	2355 2355 278( 278( 229) 223 290 217 296
ACGAAATAAAAAATAAAAAAATTCAACCAAACTCCAAAAATAAAAAA	CACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCACTGGCCCTGGA  [11]	ATCCCTACTAAACCCCACTTCCCCGACTTAAAAAACTAACCCGGAGGAGCAGTATCCGACGGAGCGGGAGCAGTATCCGACTGCGAACGGAACGGGAGCAGTATCCGACTGAACGGAACGGCACGCAC	GGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCCGCTGTGCGTCCCTTG  AAAAAATTACATGTACGCCCCCACTTTTGGGAGTTCCACCCGCTGTGCGTCCCTTG  AAAAAATTACATTAC
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ine present invention transcolated genes. The DNA sequences of signal transduction associated genes. The DNA sequences are chemically modified using a solution of bisulphite, hydrogen sulphite or disulphite. Also disclosed are oligonuclectides and/or PNA oligomers for detecting the cytosine methylation state (cpG islands) of these genes, and a method for the diagnosis and/or therapy of genetic and epigenetic parameters of genes associated with signal transduction. The genomic DNA can be obtained from cellular components which contain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certain blood, brain, heart, prostate, lung, breast or liver, histologic object slides, and all their possible combinations. The sequences of the invention are useful for the diagnosis and therapy of diseases associated with signal transduction e.g. solid tumours and sequences of different genes associated with signal transduction, or the contain of the contain the contain the contain of the contain the contain the contain of the contain the contai
                                                                                                                                                                                                                                                     Human, signal transduction associated gene; cytosine methylation state; CpG island; signal transduction associated disease; solid tumour; cancer; antitumour; cytostatic; mutant; ds.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          form part of the printed
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          Oligonucleotide for diagnosis and therapy of diseases associated with signal transduction e.g. cancer, comprises chemically modified genomic sequences of genes associated with signal transduction
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                   their complementary sequences. Note: The sequence data for this patent did not form part of the prints specification, but was obtained in electronic format directly from the European Patent Office.
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                                                                                                                                                                                                            gene modified complementary DNA #194.
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01-SEP-2000; 2000DE-1043826.
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Matches 2838; Conservative
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                                                                                                                                                                             s sequence was isolated by screening a rat ventral prostate bda gtll library in E.coli Y1090. Initial screening. With probes designed for homology to nucleotide sequences in the binding domain of known staroid receptors. Positive clones were a screened with 24mer probes specific for the various staroid eptors to eliminate those which coded for known receptors. Any ainling clones were analysed by restriction mapping and were
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/*tag=
/*toduct= full-length (902 amino acids) rAx
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11; 2428 2451 2617 2188 2248 2308 2368 2497 2557 2011 CCGGCGCCAGTTTGCTGCTGCTGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC 2128 GTGTGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTCGGGGGGATTGCATGTACG 2677 GCAAAGGTTCTCTGCTAGACGACAGCGCAGGCAAGAGCACTGAAGATACTGCTGAGTATT 2797 TCTACCCTCGGCCGCCGTCCAAGACCTACCGAGGAGCTTTCCAGAATCTGTTCCAGAGCG 1951 468 768 528 588 648 708 228 288 348 408 1891 120 Gaps 9 AGCAGCAGCAGCAGCAACAGCAGCAGGAGGTAATATCCGAAGGCAGCAGCAGCAGGAG GAGCGAGGGAGCCTCGGGGGCTCCCACTTCCTCCAAGGACAATTACTTAGGGGGCACTT 649 GAGCAAGGGACCACTGGGGGCTCCCTTCCTCCAAGGATAGTTACCTAGGGGGCAATT CGACCATTICIGACAACGCCAAGGAGTIGIGAAGGCAGIGICGGIGICCAIGGGCCTGG AGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAAGAGACTAGCCCCA---GGCAGC CTGAGAGCGGCTGCCTCCCGGAGCTGCTGCTGCTGCTCCTGGCAGGGCTCCTGCCGC AGCAGCTGCCAGCACCTCCGGACGAGGATGACTCAGCTGCCCCATCCACGTTGTCCCTGC TGGGCCCCCACTTTCCCCCGGCTTAAGCAGCTGCTCCGCTGACCTTAAAGACATCCTGAGCG |||||| ||||||||||||||||| ||| ||| AGGCCGGCACCAGCAGCAGCAGCAGCAGCAGCAGC TCTACCCACGGCCCCCGTCCAAGACCTATCGAGGAGCGTTCCAGAATCTGTTCCAGGGG AGCAGCAGCAGCAGGTGAGGATGGTTCTCCCCAAGCCCCATCGTAGAGGCCCCAAGGCT CCGAGAGAGGTTGCGTCCCAGAGCCTGGAGCCGCCGTGGCCGCCAGGAAGGGGCTGCCGC GAAGTAGGTGGAAGATTCAGCCAAGCTCAAGGATGGAAGTGCAGTTAGGGCTGGGAAGGG 191; Indels Sequence 3217 BP; 776 A; 873 C; 843 G; 725 T; 0 other; is given here. DB 12; 393; Score 2189.2; Pred. No. 0; 0; Mismatches rat AR coding sequence 0; AGGCCAGCACCATGCAACTCCTT-43.1%; 82.4%; 181 CCGGTGCCTGTTT-----Conservative Best\_Local Similarity Matches 2726; Conserv The sequenced. Query Match 349 589 2498 2618 169 2678 2738 2072 2189 2249 2309 409 2369 469 2429 529 709 829 1952 2129 289 2012 1832 1892 61 121 194 229

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            AATGTCAGCCCATCTTTCTGAATGTCCTGGAAGCCATTGAGCCAGGTGTAGTGTGTGCTG
                    GACACGACAACAACCAGCCCGACTCTTTGCAGCCTTGCTCTTAGCCTCAATGAACTGG
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                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     The sequence is used to express the corresp. peptides and for hybridisation assays of RNA and DNA encoding androgen receptors. The 98 kD product starts at the first Met codon; the 79 kD product starts from the second.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                               New DNA encoding new androgen receptor and TR2 polypeptide(s) - abito bind DNA, and derived antibodies, useful for receptor assay and
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Arag a /*tag a /*product-98 kD polypeptide
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                                                                                                                                                                                                                                                                                                                                                                                                                                                              Androgen receptor; TR2 polypeptide;
                                                                                                                                                                                                                                                                                                                                                                                                                            Rat androgen receptor DNA clone.
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                                       ACGTGGACGACCAGATGGCTGTCATTCAGTACTCCTGGATGGGGGCTCATGGTGTTTGCCA
                              TGGGCTGGCGATCCTTCACCAATGTCAACTCCAGGATGCTCTACTTCGCCCTGATCTGG
                                                           TITICAATGAGTACCGCATGCACAAGTCCCGGATGTACAGCCAGTGTGTCCGAATGAGGC
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The invention relates to a nucleic acid comprising a sequence (I) of at least 18 bases in length of a segment of chemically pre-treated genomic complement. Also included are an oligonuclectide or peptide nucleic complement. Also included are an oligonuclectide or peptide nucleic complement. Also included are an oligonuclectide or peptide incleic acid (or set thereof) of at least 9 uncleotides which hybridises to (I), probes for detecting cytosine methylation or single-complements for (I), probes for analysing diseases associated with the nucleic acids for analysing diseases associated with the complements of the CpG dinuclectides of (I). The array is useful differentiation, grading, staging, treatment and/or diagnosis of astrocytomas, or the predisposition to astrocytomas by analysing cytosine (I) astrocytomas, or the predisposition to astrocytomas by analysing cytosine by, extracting the genomic DNA, converting cytosine bases, which are numerhylated at the 5-position, in the genomic DNA sample, to uracil or another base which is dissimilar to cytosine in terms of hybridisation behaviour, by chemical treatment and amplifying chemically pre-treated genomic DNA fragments using the array and a polymerase, where the amplificates carry a detectable label. The method further, ivvolves identifying methylation status of one or more cytosine positions, and analysing methylation status of one or more cytosine positions, and cone or more data sets. The genomic DNA is chemically treated by using a cidentifying methylation status of the cytosine positions by reference to bisulphite, hydrogen sulphite or disulphite. The amplificates carry a detectable molecule fragments having a Lypical mass complements are detectable molecule fragments having a Lypical mass corrected in a mass spectrometer. The fargments of the medically have a sinnle cone-created in a mass spectrometer of pre-treated genomic nNA to have a sinnle cone-
                                                                                                                                                                  Human; ds; astrocytoma; cytostatic; staging; cysteine methyiation; CpG;
bisulphite; brain tissue; MALDI; ESI; electron spray mass spectrometry;
matrix assisted laser desorption/ionization mass spectrometry.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            Novel chemically modified genomic DNA sequences, useful in the characterisation, classification, differentiation, grading, staging; treatment and/or diagnosis of astrocytomas or predisposition to
                                                                                                                          Human DNA for staging of Astrocytomas #50.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  Claim 1; SEQ ID No 99; 37pp; English.
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    DNA; 3715
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2000DE-1043826
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ABK34012 standard;
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01-SEP-2000;
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                                        ABK34012;
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pre-treated genomic DNA to be amplified, have a single positive or negative charge for a better detectability in the mass spectrometer. Preferably, the amplificates or fragments of the amplificates are detected by matrix assisted laser desorption/ionization mass spectrometry (MALDI) or using electron spray mass spectrometry (ESI). The

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                                                             Length 3715;
             patent did not form part
was obtained in electronic
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                                                 other;
 pre-treated
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                                                            Score 2154; DB
Pred. No. 0;
0; Mismatches
        samples of the invention.

Note: The sequence data for this patent did
Note: the printed specification, but was obtain
format directly from WIPO at
 chemically
                                    ftp.wipo.int/pub/published_pct_sequences
                                                841 A; 149 C; 1001
  sequence is one of the
                                                               42.48;
                                                                      Similarity 74.5
32; Conservative
                                                 Sequence 3715 BP;
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                  CTGATGTGTGCTACCCTGGCGGCATGGTGAGAGAGTGCCCTATCCCAGTCCCACTTGTG
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Human; signal transduction associated gene; cytosine methylation state; CpG island; signal transduction associated disease; solid tumour; cancer; antitumour; cytostatic; mutant; ds.
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3044 ATCGTATTATTGTAAAAAAAAAAATTTTATATTTTGTTTAAGACGTTTTATTAGT 3103
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                                                                                                                     TCTCTGTGCAAAGTGCCCAAGATCCTTTCTGGGAAAGTCAAGCCCCATCTATTTCCACACCC
                                                                                                                                                                                      TCACCAAGCTCCTGGACTCCGTGCAGCCTATTGCGAGAGAGCTGCATCAGTTTTTG
                                                                                                                                                                                                                                     AGTGAAGCATTGGAAACCCTATTTCCCCACCCCAGCTCATGCCCCCTTTCAGATGTCTTC
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The present invention relates to chemically modified DNA sequences of signal transduction associated genes. The DNA sequences are chemically modified using a solution of bisulphite, hydrogen sulphite or calculphite. Also disclosed are oligonuclectides and/or PNA oligomers for detecting the cytosine methylation state (CpG islands) of these genes, and a method for the diagnosis and/or therapy of genetic and cepigenetic parameters of genes associated with signal transduction. The genomic DNA can be obtained from cells or cellular components which contain DNA, e.g. cell lines, biopsies, blood, sputum, stool, urine, certeal-spinal fulmi, tissue embedded in paraffin such as tissue from eyes, intestine, kidney, brain, heart, prostate, lung, breast or liver, histologic object slides, and all their possible combinations. The sequences of the invention are useful for the diagnosis and therapy of diseases associated with signal transduction, or sequences of different genes associated with signal transduction, or their complementary sequences.

Sequence of different genes associated with signal transduction, or their complementary sequences.

Oute: The sequence data for this patent did not form part of the printed process. Oligonucleotide for diagnosis and therapy of diseases associated with signal transduction e.g. cancer, comprises chemically modified genomic sequences of genes associated with signal transduction . Claim 1; SEQ ID No 387; 24pp; English χ. Piepenbrock C, European Patent Office. WPI; 2002-147896/19 olek A, 

Sequence 3715 BP; 841 A; 149 C; 1001 G; 1724 T; 0 other;

1639 1699 1700 CITITIGGGIGGIIGCICCCCCCAAGIIICCIICIIGGAGCIICCCGCAGGIGGGCAGCI 1759 GGCGGGGTAAGGGAAGTGCGAGATTCAGCCAAGCTCAAGGATGGAAGTGCAGTTAG 1879 1340 GCGGAGAGAACCCTCTGTTTTCCCCCCACTCTCTCCACCTCCTCCTGCCTTCCCCACCC 1399 1459 AGCTGCAGCGACTACCGCATCATCACAGCCTGTTGAACTCTTCTGAGCAAGAGAAGGGGA 1819 367 427 187 89 CGAGTGCGGAGCCAGAGATCAAAAGATGAAAAGGCAGTCAGGTCTTCAGTAGCCAAAAAA AGGGCAGATCTTGTCCACCGTGTGTCTTCTTGTGCAGAGCTTTGAGGCTGTCAGAGCG TITITIGCGTGGTTGTTTTCGTAAGTTTTTTTTTTTGGAGTTTTTCGTAGGTGGGTAGTT 45; Query Match
Best Local Similarity 74.5%; Pred. No. 0;
Matches 2782; Conservative 0; Mismatches 905; Indels 45; 1580 1640 09/1 1820 1520 188 248 308 368 428 1400 δλ Q g ò 셤 g δ g δ q δŽ g οy ò ò ò

2659 2779 2839 2239 2059 883 607 667 AAGATACTGCTGAGTATTCCCCTTTCAAGGGAGGTTACACCAAAGGGCTAGAAGGCGAGA GCCTAGGCTGCTCTGGCAGCGCTGCAGCAGGAGCTCCGGGACACTTGAACTGCGGTCTA GTTTAGGTTCTTTTGGTAGCGTTGTAGTAGGGAGTTTCGGGATATTTGAATTGTCGTTTA CCCTGTCTCTCTACAAGTCCGGAGCACTGGACGAGGCAGCTGCGTACCAGAGTCGCGACT TGTCCCTGCTGGGCCCCCACTTTCCCCGGCTTAAGCAGCTGCTCCGCTGACCTTAAAGACA GTGCCCCATTGGCCGAATGCAAAGGTTCTCTGCTAGACGACAGCGCAGGCAAGAACACTG 2120 CCAGGCAGCAGCAGCAGCAGCAGGGTGATGGTTCTCCCCCAAGCCCATCGTAGAGGCC CCACAGGCTACCTGGTCCTGGATGAGGAACAGCAACCTTCACAGCCGCAGTCGGCCCTGG 2240 AGTGCCACCCGAGAGAGGTTGCGTCCCAGAGCCTGGAGCCGCCGTGGCCGCCAGCAAGG ATTACTTAGGGGGCCACTTCGACCATTTCTGACAACGCCAAGGAGTTGTGTAAGGCAGTGT CGGTGTCCATGGGCCTGGGTGTGGAGGCGTTGGAGCATCTGAGTCCAGGGGAACAGCTTC GGGGGGATTGCATGTACGCCCCACTTTTGGGAGTTCCACCCGCTGTGCGTCCCACCTT GGCGGGGTAAGGGAAGTAGGTGGAAGATTTAGTTTAAGTTTAAGGATGGAAGTGTAGTTAG 764 TTAGGTAGTAGTAGTAGTAGGGGGGGGGGTTTTTTTTAAGTTTATCGTAGAGGTT TCCTGAGCGAGGCCAGCACCATGCAACTCCTTCAGCAACAGCAGCAGGAAGCAGTATCCG AAGGCAGCAGCAGGGAGAGCGAGGGAGGCCTCGGGGGGTCCCACTTCCTCCAAGGACA GGCTGGGAAGGGTCTACCCTCGGCCGCTCCAAGACCTACCGAGGAGCTTCCAGAATC GCGTAGTATTTTCGGCGTTAGTTTGTTGTTGTT---------GT TGTTCCAGAGCGTGCGCGAAGTGATCCAGAACCCGGGCCCCCAGGCCACCCAGAGGCCGGA 2840 2780 1484 2300 2360 2420 1064 2480 1124 2540 1184 2600 2660 2720 2060 2180 1880 1940 2000 884 548 809 899 704 g qq g ŏ g δ g ŏ g ö q g ŏ ŏ δ g QΥ g ŏ g δy g Qγ g ò g δ δ g g

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4210 4270 4150 2923 TCACCCCCAGGAATICCIGIGCAIGAAAGCACIGCIACICIICAGCAFIATICCAGIGG 4330 ATGGGCTGAAAAATCAAAAATTCTTTGATGAACTTCGAATGAACTACATCAAGGAACTCG 4390 4451 TCACCAAGCICCTGGACTCCGTGCAGCCTATTGCGAGAGAGAGCTGCATCAGTTTAG 4510 4511 ACCTGCTAATCAAGTCACATGGTGAGCGTGGACTTTCCGGAAATGATGGCAGAGTCA 4570 4450 3164 ATTTGTTAATTAAGTTATATATGTGAGGTGGATTTTTCGGAAATGATGGTAGATTA 3223 4571 TCTCTGTGCAAGTGCCCAAGATCCTTTCTGGGAAAGTCAAGCCCATGTATTTCCACACC 4630 4690 3343 TGCCTGTTATAACTCTGCACTACTCCTCTGCAGTGCCTTGGGGAATTTCCTCTATTGATG 4750 4870 3512 4930 3572 4810 ACTCCTGGATGGGGCTCATGGTGTTTGCCATGGGCTGGCGATCCTTCACCAATGTCAACT CCAGGATGCTCTACTTCGCCCCTGATCTGGTTTTCAATGAGTACCGCATGCACAAGTCCC GGATGTACAGCCAGTGTGTCCGAATGAGGCACCTCTCTCAAGAGTTTGGATGGCTCCAAA ATCGTATCATTGCATGCAAAAAAAAATCCCACATCCTGCTCAAGACGCTTCTACCAGC AGTGAAGCATTGGAAACCCTATTTCCCCAACCTCATGCCCCCTTTCAGATGTCTTC CICCIIICIIIIICIICCIICCCICCCIAICIAACCCICCCAIGGCACCIILAGACIIIIGC TCAAACAAAA 5062 4151 4091 2804 2864 4211 4271 2924 4331 4391 4631 4691 3344 3453 4871 4751 4811 3513 4991 3633 5051 òγ οp Ω Ω οy q ΟŽ Db Qy qq Op δ δy Db QΥ q QQ οy Qγ Ωp Ωÿ QQ Oy Op  $_{2}^{0}$ QQ QΥ qq QΥ QQ QΥ QQ ŏλ Dp οy

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                                                                                                                                                                                                                                                                                                                                 The present invention relates to a method for analysis of inactivation of x chromosome. The method comprises analysing methylation of human androgen receptor (HUMARA) gene (the present sequence) by a methyl-specific polymerase chain reaction (PCR). The PCR primers amplify the base sequence of the region containing repeated number of polymorphism of CAG of the HUMARA gene, specific to methylation of the cytosine base at the 199th, the 203rd and the 206th or the 296th position of the present sequence. The method is useful for the detection of uniformity of cell growth, that is clonality.
                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                1940
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                                                                                            androgen
                                                                                                                                                                                                                                                                                                                                                                                                                                                            33.8%; Score 1715.6; DB 2:
98.7%; Pred. No. 1.8e-284;
iive 0; Mismatches 4;
                                                                                            Human; PCR primer; X chromosome inactivation; methylation; HUMARA; ds.
                                                                                                                                                                                                                                                                                                                    Claim 2; Page 11-12; 14pp; Japanese.
                 ВÞ
                                                                                                                                                                                                                             (MITP ) MITSUBISHI YUKA BCL KK.
                                                                            Human androgen receptor gene
                  DNA; 1810
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Matches 1767; Conservative
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                 AAF84342 standard;
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                                                                                                                              Homo sapiens
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                                   CAGGCAGCAGCAGCAGCAGCAGGGTGAGGATGGTTCTCCCCCAAGCCCATCGTAGAGGCCC
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